



UP-MEDICINE SPECIALTY CLINIC

Patient Name: [REDACTED]
MRN: [REDACTED]
Financial #: [REDACTED]
Admit Date: 11/8/2022
Discharge Date: 11/8/2022
Patient Type: Clinic
DOB/Age/Sex: [REDACTED] Male
Attending: Dandachi MD,Dima
Referring: Referred,Self

Clinic Notes

DOCUMENT NAME:
RESULT STATUS:
SIGN INFORMATION:

Infectious Disease IM Clinic Note
Modified
Dandachi MD,Dima (11/15/2022 11:41 CST); Dandachi MD,
Dima (11/15/2022 11:41 CST); Escovar MD,Javier
(11/8/2022 10:46 CST)

Chief Complaint

follow up

History of Present Illness

38-year-old male coming for follow-up of HIV. Feeling well, no acute complaints at this time. Is on Biktarvy and Bactrim. 7 days missed doses of Biktarvy in the last 30 days. Zero missed doses of Bactrim. Viral load UD ,8/2022. CD4 count 266 ,8/2022. Tolerating medication well. No adverse effects. Is sexually inactive. Denies current drug use.

Review of Systems

14 point ROS negative unless otherwise specified

Physical Exam

Vitals & Measurements

HR: 71 BP: 139/86 SpO2: 98%
HT: 182 cm WT: 82.9 kg BMI: 25

Gen: Calm, cooperative. NAD.

Skin: No pathologic lesions

HEENT: EOMI, PERRLA, no rhinorrhea, conjunctival injection or suffusion

Pulm: CTA B/L. No Rhonchi, wheezes, or rales.

Cardio: RRR, Regular S1/S2, no rubs murmurs or gallops

Vasc: +2 pulses PT/DP. No edema

GI: Soft, NT, ND, BS+, no rebound or guarding

Neuro: AOX3, responds to questions appropriately

MSK: 5/5 strength U/LE B/L

Assessment/Plan

37 yr old M patient presenting to HIV clinic for follow up

HIV/AIDs subtype group B

- Year of Dx 1/2022 HIV mode of transmission: MSM
- VL UD 8/2022
- CD4 266 8/2022
- Current ART Biktarvy 5/27/2022 - P ,
- Missed 7 doses reportedly due to prison unable to obtain meds, no reported SE
- Prophylaxis: Bactrim 5/27/2022 - P
- ART genotype 5/24/2022 wild type

Problem List/Past Medical History

Ongoing

No qualifying data

Historical

No qualifying data

Medications

Bactrim DS 800 mg-160 mg oral tablet, 1
Tablet(s), Oral, Daily, 1 refills
Biktarvy 50 mg-200 mg-25 mg oral tablet, 1
Tablet(s), Oral, Daily, 6 refills
Biktarvy 50 mg-200 mg-25 mg oral tablet, 1
Tablet(s), Oral, Daily, 5 refills

Allergies

NKA

Social History

Smoking Status

Current every day smoker

Family History

Denied family history of heart disease

Immunizations

Vaccine	Date	Status
meningococcal conjugate Vaccine	08/09/2022	Given
zoster vaccine, Inactivated	08/09/2022	Given
pneumococcal 20-valent conjugate vaccine	05/24/2022	Given
meningococcal conjugate Vaccine	05/24/2022	Given

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If you have any questions or concerns regarding this documentation, please call (573) 882-3170.

Report Request ID: 453592762

Page 1 of 10

Print Date/Time: 11/21/2022 16:12 CST

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Clinic Notes

- HLA B5701 negative

Screening and health maintenance

- Hx of OI n/a
- Quantiferon negative (CD4 124)
- GC NAAT negative x3
- Syphilis screening negative
- AFB blood Cx negative
- Tdap booster pending
- Hep A nonimmune: today
- Hep B : Immune by exposure Hep B s Ab and c Ab +. s Ag negative
- Hep C negative 5/2022
- PCV 20 5/2022
- Menactra (MenACWY-DT) x2 8/2022
- HPV today
- Shingrix x2 8/2022
- Flu vaccine 11/2022, patient reported
- COVID-19 x 3

Plan

- Labs will check today: CBC, CMP, VL ,CD4, QFN (now that CD4 has recovered), Lipid panel
- Immunization: HPV, Hep A
- Continue Biktarvy and Bactrim
- Communicate results to RN (?) as office hill is on medical leave. 660-827-0056 ext 2?

2. Substance use

Previously reported Methamphetamine IV use and Cocaine
Today denies any past history of use despite reporting use last visit

3. Hx of gonorrhea s/p treatment

- Negative oral and rectal

Staffed with Dr. Dandachi

RTC pending VL, if unremarkable 3 months

Vaccine	Date	Status
zoster vaccine, inactivated	05/24/2022	Given
SARS-CoV-2 (COVID-19) mRNA BNT-162b2 vax	01/11/2022	Recorded
SARS-CoV-2 (COVID-19) mRNA BNT-162b2 vax	04/22/2021	Recorded
SARS-CoV-2 (COVID-19) mRNA BNT-162b2 vax	04/01/2021	Recorded
pneumococcal 23-valent vaccine	-	Not Given
tetanus-diphtheria toxoids	08/12/1999	Recorded
measles/mumps /rubella virus vaccine	10/04/1990	Recorded
poliovirus vaccine live trivalent	11/17/1989	Recorded
poliovirus vaccine live trivalent	01/16/1987	Recorded
DTP (old vaccine)*	12/19/1986	Recorded
poliovirus vaccine live trivalent	02/20/1986	Recorded
DTP (old vaccine)*	02/20/1986	Recorded
measles/mumps /rubella virus vaccine	10/10/1985	Recorded
poliovirus vaccine live trivalent	01/25/1985	Recorded
DTP (old vaccine)*	01/25/1985	Recorded
poliovirus vaccine live trivalent	11/23/1984	Recorded

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Report Request ID: 453592762

Page 2 of 10

Print Date/Time: 11/21/2022 16:12 CST

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Clinic Notes

Vaccine	Date	Status
DTP (old vaccine)*	11/23/1984	Recorded
poliovirus vaccine live trivalent	09/14/1984	Recorded
DTP (old vaccine)*	09/14/1984	Recorded

Visit Information

Attending Provider: Attending Default
Referring Provider: Self Referred
Original Referring Provider: Self Referred
Primary Care Provider: Deedee E Gilmore
FNP
Visit Date: 11/08/2022

Addendum by Dandachi MD, Dima on November 15, 2022 11:41:47 CST

I personally saw and evaluated the patient on the date found in the header of this document. I reviewed the history, physical, labs, micro, medications, and imaging studies. I have discussed the management of the patient with the author of this document and I agree with the findings and plan as documented above.

Dima Dandachi, MD, MPH
Assistant Professor of Clinical Medicine
Division of Infectious Diseases
University of Missouri-Columbia

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Report Request ID: 453592762

Page 3 of 10

Print Date/Time: 11/21/2022 16:12 CST

Patient Name: [REDACTED]
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Admit Date: 11/8/2022
Discharge Date: 11/8/2022

General Lab Results

Hematology

Collected Date 11/8/2022
Collected Time 10:24 CST

Procedure		Units	Reference Range
WBC	4.76 ¹¹	x10(9)/L	[3.50-10.50]
RBC	4.46 ¹¹	x10(12)/L	[4.32-5.72]
HGB	14.0 ¹¹	g/dL	[13.5-17.5]
HCT	41.2 ¹¹	%	[38.8-50.0]
MCV	92.4 ¹¹	fL	[81.2-95.1]
MCH	31.4 ¹¹	pg	[26.0-33.0]
MCHC	34.0 ¹¹	g/dL	[32.0-36.0]
RDW SD	46.1 ¹¹	fL	[35.1-43.9]
RDW CV	13.6 ¹¹	%	[11.8-15.6]
PLT	223 ¹¹	x10(9)/L	[150-450]
MPV	10.1 ¹¹		[8.0-12.0]
% Neutrophils	43.2 ⁰¹¹¹	%	
Absolute Granulocytes	2.05 ⁰¹¹¹	x10(9)/L	[1.70-7.00]
% Immature Granulocytes	.40 ⁰¹¹¹¹¹	%	[0.02-0.42]
Abs Immature Granulocytes	0.02 ⁰¹¹¹¹¹	x10(9)/L	[0.00-0.03]
% Lymphocytes	36.3 ⁰¹¹¹	%	
Abs Lymphocytes	1.73 ⁰¹¹¹	x10(9)/L	[0.90-2.90]
% Monocytes	13.0 ⁰¹¹¹	%	
Abs Monocytes	0.62 ⁰¹¹¹	x10(9)/L	[0.30-0.90]
% Eosinophils	6.7 ⁰¹¹¹	%	
Abs Eosinophils	0.32 ⁰¹¹¹	x10(9)/L	[0.05-0.50]
% Basophils	0.4 ⁰¹¹¹	%	
Abs Basophils	0.02 ⁰¹¹¹	x10(9)/L	[0.00-0.30]
% Nucleated RBCs	0.0 ¹¹	%	
Absolute Nucleated RBCs	0.0 ¹²¹¹	x10(9)/L	[0.0-0.0]

Order Comments

O1: Auto differential (Auto diff)
Added by Discern rule.

Interpretive Data

i1: % Immature Granulocytes, Abs Immature Granulocytes

Add'l Patient Type	%IG	IG 10^9/L
Pregnancy: 1st Trimester	0.04 - 0.92	0.003 - 0.091
2nd Trimester	0.10 - 2.00	0.007 - 0.247
3rd Trimester	0.20 - 3.80	0.018 - 0.456

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Report Request ID: 453592762

Page 4 of 10

Print Date/Time: 11/21/2022 16:12 CST

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General Lab Results

Hematology

Interpretive Data

i2: Absolute Nucleated RBCs
Normal values not established in patients less than 18 years old.

Chemistry

Collected Date	11/8/2022		
Collected Time	10:24 CST		
Procedure		Units	Reference Range
Sodium	140 ¹¹	mmol/L	[136-145]
Potassium	3.8 ¹¹	mmol/L	[3.5-5.1]
Chloride	105 ¹¹	mmol/L	[98-107]
CO2	25 ¹¹	mmol/L	[22-29]
Anion gap	14 ¹¹	mmol/L	[0-20]
Glucose Lvl	87 ¹¹	mg/dL	[70-139]
BUN	15 ¹¹	mg/dL	[6-20]
Creatinine, standardized	1.20 ¹¹	mg/dL	[0.70-1.20]
Estimated GFR for Adults	79 ¹¹ ^{R113}	mL/min/1.73m ²	[>=90]
Estimated GFR for peds	Not calculated ¹⁴	mL/min/1.73m ²	[>=90.00]
Calcium	9.1 ¹¹	mg/dL	[8.6-10.2]
Total Protein	7.8 ¹¹	g/dL	[6.6-8.7]
Albumin	4.6 ¹¹	g/dL	[3.5-5.2]
T Billi	0.21 ¹¹	mg/dL	[0.00-1.60]
Alkaline Phosphatase	92 ¹⁵	units/L	[40-129]
AST-SGOT	23 ¹¹	units/L	[<=40]
ALT-SGPT	28 ¹¹	units/L	[10-50]
Cholesterol	157 ¹¹	mg/dL	[0-200]
HDL Cholesterol	38 ¹¹ ^{L16}	mg/dL	[40-60]
Cholesterol HDL Ratio	4.1 ¹⁷		[<=4.9]
LDL (Calculated)	92 ¹⁰	mg/dL	[0-129]
Triglycerides	135 ¹⁰	mg/dL	[0-150]
(M TB) NIL value	0.527 ¹²	IU/mL	
(M TB) Mitogen	>10.000 ¹²	IU/mL	
(M TB) Antigen1-NIL value	-0.039 ¹²	IU/mL	
(M TB) Antigen2-NIL Value	-0.043 ¹²	IU/mL	
(M TB) Mitogen-NIL value	>9.473 ¹²	IU/mL	
(M TB) QFT-Plus by EIA	Negative ¹¹⁰		
TB1 Antigen	0.488 ¹²	IU/mL	
TB2 Antigen	0.484 ¹²	IU/mL	

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Report Request ID: 453592762

Page 5 of 10

Print Date/Time: 11/21/2022 16:12 CST

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Discharge Date: 11/8/2022

General Lab Results

Chemistry

Result Comments

R1: Estimated GFR for Adults

The eGFR was estimated using the CKD-EPI equation. The National Kidney Foundation recommends that all clinical labs utilize this equation: Levey AS, Stevens LA, Schmid CH, et al. A new equation to estimate glomerular filtration rate. Ann Intern Med. 2009;150(9):604-612.

For calculation reference see https://www.kidney.org/professionals/kdoqi/gfr_calculator

Interpretive Data

i3: Estimated GFR for Adults

Changed to CKD-EPI 2021 on Nov. 18th, 2021.

i4: Estimated GFR for peds

The estimated GFR was calculated using the "Bedside Schwartz equation (2009)".

Reference: Pediatric GFR calculator at National Kidney Foundation Website.

i5: Alkaline Phosphatase

Normal range for plasma alkaline phosphatase in females 20 years or older:

35-104 U/L according to manufacturer's instructions

35-129 U/L according to data analysis on adult females who presented to MUHC in 2022 without a significant diagnosis

i6: Clinical judgement is advised.

HDL Cholesterol

HDL Cholesterol Level mg/dL	Category
-----------------------------	----------

Less than 40(for Men)	Low HDL cholesterol. A major risk factor for heart disease.
Less than 50 (for Women)	

60 and above	High HDL cholesterol. An HDL of 60 mg/dL and above is considered protective against heart disease.
--------------	--

National Cholesterol Education Program

NHLBI Health Information Network

P.O. Box 30105

Bethesda MD 20824-0105

<http://www.nhlbi.nih.gov>

i7: Cholesterol HDL Ratio

The cholesterol HDL ratio is the best overall predictor of heart disease.

-Less than 5 is normal

-From 5 to 9 there is increasing risk

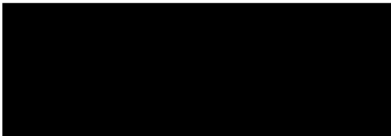

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Page 6 of 10

Print Date/Time: 11/21/2022 16:12 CST

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General Lab Results

Chemistry

Interpretive Data

i7: Cholesterol HDL Ratio

-Higher than 9 is high risk. A ratio higher than 9 will often require prescription medication to help prevent heart disease.

See <http://fcm-algo.umh.edu/Algorithms/Lipid.htm> for more information.

i8: LDL (Calculated)

LDL Cholesterol Level mg/dL	Category
-----------------------------	----------

Less than 100	Optimal
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100 to 129	Near or above optimal
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130 to 159	Borderline high
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160 to 189	High
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190 and above	Very High
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Note: "Values < 80 mg/dL may indicate hypobetalipoproteinemia, if not on Statin therapy."

Reference: ATP III Guidelines, <http://fcm-algo.umh.edu/Algorithms/Lipid.htm>
National Cholesterol Education Program
NHLBI Health Information Network
P.O. Box 30105
Bethesda MD 20824-0105
<http://www.nhlbi.nih.gov>

Footnote: Depending on cardiac risk factors, age, and sex, UHC Cardiology prefers:

<75 mg/dl optimal
<100 mg/dl desirable

i9: Triglycerides

Less than 150	Normal
---------------	--------

150 - 199	Borderline high
-----------	-----------------

200 - 499	High
-----------	------

500 and above	Very high
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Report Request ID: 453592762

Page 7 of 10

Print Date/Time: 11/21/2022 16:12 CST

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Discharge Date: 11/8/2022

General Lab Results

Chemistry

Interpretive Data

i9: Triglycerides

National Cholesterol Education Program
NHLBI Health Information Network
P.O. Box 30105
Bethesda, MD 20824 - 0105
<http://www.nhlbi.nih.gov>

i10: (M TB) QFT-Plus by EIA
QFT-Plus is an indirect test for M. tuberculosis infection.

See general guidance on the diagnosis and treatment of TB disease and LTBI
(<https://www.cdc.gov/tb/publications/guidelines/default.htm>).

Mycobacterium Tuberculosis by ELISA results are interpreted using the following criteria:

Nil (IU/mL)	TB1 minus Nil	TB2 minus Nil	Mitogen minus Nil (IU/mL)	QFT-Plus Result	Interpretation
<=8.0	>= 0.35 and >= 25% of Nil	Any	Any	Positive	M. tuberculosis infection likely
<=8.0	Any	>= 0.35 and >= 25% of Nil	Any	Positive	M. tuberculosis infection likely
<=8.0	< 0.35 or >= 0.35 and >= 25% of Nil	< 0.35 or >= 0.35 and >= 25% of Nil	>=0.50	Negative	M. tuberculosis infection NOT likely
<=8.0	< 0.35 or >= 0.35 and >= 25% of Nil	< 0.35 or >= 0.35 and >= 25% of Nil	<0.50	Indeterminate	Likelihood of M. tuberculosis infection cannot be determined
>8.0	Any	Any	Any	Indeterminate	Likelihood of M. tuberculosis infection cannot be determined

Reference Range: (Negative) Non-Responsive to ESAT-6, CFP-10 and/or TB7.7 antigens.

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Report Request ID: 453592762

Page 8 of 10

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General Lab Results

Chemistry

Interpretive Data

i10: (M TB) QFT-Plus by EIA

Results from QFT-Plus testing must be used in conjunction with each individual's epidemiological history, current medical status, and other diagnostic evaluations.

The performance of the USA format of the QFT-Plus test has not been extensively evaluated with specimens from the following groups of individuals:

- Pregnant women
- Individuals younger than age 17 years
- Individuals who have impaired or altered immune functions.

Performed on DiaSorin Liaison XL.

Molecular Pathology

Collected Date 11/8/2022

Collected Time 10:24 CST

Procedure	Units	Reference Range
HIV-1 Quant	<30 ^{*2} copies/mL	
HIV-1 Log10	<1.47 ^{*2} log10	
HIV-1 Reflex Testing Ordered	No ^{*2}	

Performing Locations

- *1: This test was performed at:
University of Missouri Health Care, University Hospital, Laboratory, 1 Hospital Dr., Columbia, MO, 65212- , US,
573-882-1400
- *2: This test was performed at:
APG Lab, A. P. Green Bldg, 201 Business Loop 70W, Columbia, MO, 65203- , US

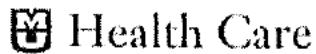
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Report Request ID: 453592762

Page 9 of 10

Print Date/Time: 11/21/2022 16:12 CST



UP-MEDICINE SPECIALTY CLINIC

Patient Name:

MRN:

Financial #:

Admit Date: 11/8/2022

Discharge Date: 11/8/2022

LAB- Flow Cytometry

Collected Date	11/8/2022
Procedure	
% CD3 T Cells	68.6 ^{*1}
% CD4 T Helper Cells	13.4 ^{L*1}
% CD8 T Suppressor Cells	50.6 ^{H*1}
CD3 T Cells	1287.0 ^{*1}
CD4 T Helper Cells	250.0 ^{L*1}
CD8 T Suppressor Cells	949.00 ^{H*1}
CD4/CD8 Ratio (Help/Suppr)	0.26 ^{L*1}
Flow Cytometry Compliance Comment	See Comment ^{111*1}

Interpretive Data

i11: Flow Cytometry Compliance Comment

Analyte Specific Reagent: This test was developed and its performance characteristics determined by the Clinical Flow Cytometry Laboratory, UMHC-Integrated Central Pathology Labs, University of Missouri Health Care. It has not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance is not necessary. This test is used for clinical purposes, and should not be regarded as investigational or for research. This laboratory is certified under the Clinical Laboratory Improvement Amendment of 1988 (CLIA-88) as qualified to perform high complexity clinical laboratory testing.

Performing Locations

*1: This test was performed at:

University of Missouri Health Care, University Hospital, Laboratory, 1 Hospital Dr., Columbia, MO, 65212- , US, 573-882-1400

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Page 10 of 10

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